Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

Claim1 (currently amended): A plant or yeast eukaryotic cell that comprises a prokaryotic recombinase polypeptide or a nucleic acid that encodes the prokaryotic recombinase, wherein the recombinase is capable of mediating site-specific recombination in the eukaryotic cell between an *attB* recombination site and an *attP* recombination site to form an *attL* and an *attR* site; and wherein the recombinase is not capable of mediating in the eukaryotic cell recombination between the *attL* site and the *attR* site, wherein the recombinase is a bacteriophage ΦC31 integrase; and only one *attP* or only one *attB* recombination site of bacteriophage ΦC31 integrase integrated in its genome.

Claims 2 to 5 (canceled).

Claim 6 (previously presented): The eukaryotic cell of claim 1, wherein the cell comprises a nucleic acid that comprises a coding sequence for the recombinase polypeptide, which coding sequence is operably linked to a promoter that mediates expression of the recombinase-encoding polynucleotide in the eukaryotic cell.

Claim 7 (original): The eukaryotic cell of claim 6, wherein the nucleic acid further comprises a selectable marker.

Claim 8 (original): The eukaryotic cell of claim 6, wherein the promoter is an inducible or a repressible promoter.

Claim 9 (canceled).

Claim 10 (previously presented): The eukaryotic cell of claim 1, wherein the cell is a yeast cell.

Claim 11 (previously presented): The eukaryotic cell of claim 1, wherein the eukaryotic cell is a plant cell.

Claims 12 to 35. (canceled).

Claim 36 (currently amended): A plant or yeast eukaryotic cell that comprises: an attP or attB recombination site of bacteriophage Φ C31 integrase integrated in its genome; and

a non-genomic nucleic acid comprising a heterologous nucleic acid or a transgene, and an **only one** attP site **of bacteriophage** Φ **C31 integrase** if the cell has the genomic attB site or **only one** an attB site **of bacteriophage** Φ **C31 integrase** if the cell has the genomic attP site; wherein the eukaryotic cell further comprises a Φ C31 integrase polypeptide.

Claim 37 (previously presented). The eukaryotic cell of claim 36, wherein the non-genomic nucleic acid comprises the transgene.

Claims 38 to 42 (canceled).

Claim 43 (previously presented): The eukaryotic cell of claim 36, wherein the eukaryotic cell comprises a nucleic acid that comprises a polynucleotide that encodes the Φ C31 integrase polypeptide.

Claim 44 (original). The eukaryotic cell of claim 43, wherein the nucleic acid further comprises a selectable marker.

Claim 45 (previously presented): The eukaryotic cell of claim 43, wherein the nucleic acid further comprises an inducible promoter which controls expression of the Φ C31 integrase-encoding polynucleotide in the cell.

Claim 46 (canceled).

Claim 47 (previously presented): The eukaryotic cell of claim 36, wherein the plant is a dicot or a monocot.

Claims 48 to 51 (canceled).

Claim 52 (currently amended): A eucaryotic somatic cell in culture comprising: a prokaryotic recombinase polypeptide or a nucleic acid that encodes the prokaryotic recombinase, wherein the recombinase is capable of mediating site-specific recombination in the eukaryotic cell between an *attB* recombination site and an *attP* recombination site to form an *attL* and an *attR* site, and is not capable of mediating in the eukaryotic cell recombination between the *attL* site and the *attR* site;

the attP or attB recombination site integrated in its genome;

a non-genomic nucleic acid comprising a transgene or a heterologous nucleic acid and **an** <u>only one</u> <u>attP</u> site if the cell has the genomic <u>attB</u> site or <u>only one</u> an <u>attP</u> site if the cell has the genomic <u>attB</u> site;

wherein the recombinase is a bacteriophage ΦC31 integrase and the attP and attB sites are bacteriophage ΦC31 integrase recombination sites.

Claims 53 to 60 (canceled).

Claim 61 (currently amended): A method for obtaining site-specific recombination in a eukaryotic cell, the method comprising:

providing a eukaryotic cell that comprises an *attB* recombination site <u>and or</u> an *attP* recombination site <u>of bacteriophage Φ C31 integrase integrated in its genome and a non-genomic nucleic acid comprising a transgene or a heterologous nucleic acid and only one *attP* site if the cell has the genomic *attB* site or only one *attP* site if the cell has the genomic *attB* site;</u>

contacting the attB and the attP recombination sites with a prokaryotic recombinase polypeptide, resulting in recombination between the recombination sites, thereby forming an attR and an attL recombination site;

wherein the recombinase polypeptide can mediate site-specific recombination between the *attB* and *attP* recombination sites, but cannot mediate recombination between the *attR* and *attL* recombination sites;

wherein the recombinase is-a bacteriophage Φ C31 integrase,

Claim 62 (previously presented) The method of claim 61, wherein the eukaryotic cell is selected from the group consisting of a yeast cell, a fungal cell, a plant cell, an insect cell and an animal cell.

Claims 63 to 64 (canceled).

Claim 65 (currently amended): The method <u>-of-claim 61 for obtaining site-specific recombination in a eukaryotic cell, the method comprising:</u>

providing a eukaryotic cell that comprises an attB recombination site and-an attP recombination site of bacteriophage Φ C31 integrase, wherein the attB recombination site and the attP recombination site are present on a single nucleic acid molecule;

contacting the attB and the attP recombination sites with a prokaryotic recombinase polypeptide, resulting in recombination between the recombination sites, thereby forming an attR and an attL recombination site;

wherein the recombinase polypeptide can mediate site-specific recombination between the attB and attP recombination sites, but cannot mediate recombination between the attR and attL recombination sites; and

wherein the recombinase is-a bacteriophage ΦC31 integrase.

Claim 66 (previously presented): The method of claim 65, wherein the *attB* recombination site and the *attP* recombination site are in a direct orientation and the recombination results in excision of the portion of the nucleic acid molecule that lies between the *attB* and *attP* recombination sites.

Claim 67 (canceled)

Claim 68 (currently amended): The method of claim 65, wherein the *attB* recombination site and the *attP* recombination site are in an inverted orientation and the recombination results in inversion of the portion of the nucleic acid molecule that lies between the *attB* and *attP* recombination sites.

Claim 69 (canceled).

Claim 70 (previously presented): The method of claim 61, wherein the eukaryotic cell comprises a polynucleotide that encodes the recombinase polypeptide.

Claim 71 (new). A plant regenerated from a plant eucaryotic cell of claim 36.